

United States Patent and Trademark Office



UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 • Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/749,143	12/29/2003	W. James Jackson	069748-0304	3867
22428 7590 09/10/2007 FOLEY AND LARDNER LLP		EXAMINER		
SUITE 500 3000 K STREE	ET NW		DEVI, SARVAMANGALA J N	
WASHINGTON, DC 20007			ART UNIT	PAPER NUMBER
			1645	
·			MAIL DATE	DELIVERY MODE
			09/10/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

 -		Application No.	Applicant(s)			
Office Action Summary		10/749,143	JACKSON ET AL.			
		Examiner	Art Unit			
		S. Devi, Ph.D.	1645			
Period fo	The MAILING DATE of this communication app or Reply	ears on the cover sheet with the c	orrespondence address			
WHIC - Exter after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DANSIONS of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. O period for reply is specified above, the maximum statutory period we are to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing ed patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim rill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	I. lely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status		•				
1)🖂	Responsive to communication(s) filed on 11 Ju	ily 2007.				
2a)⊠	This action is FINAL . 2b) This action is non-final.					
3)	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.					
Dispositi	ion of Claims	•				
5)□ 6)⊠ 7)□	Claim(s) 1-7,14-37 and 52 js/are pending in the 4a) Of the above claim(s) is/are withdraw Claim(s) is/are allowed. Claim(s) 1-7, 14-37 and 52 js/are rejected. Claim(s) is/are objected to. Claim(s) are subject to restriction and/or	vn from consideration.				
Applicati	ion Papers					
10)□	The specification is objected to by the Examiner The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the Replacement drawing sheet(s) including the correction of the oath or declaration is objected to by the Examiner.	epted or b) objected to by the Edrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).			
Priority u	under 35 U.S.C. § 119					
12) a)[Acknowledgment is made of a claim for foreign All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priority application from the International Bureau See the attached detailed Office action for a list of	s have been received. s have been received in Application ity documents have been receive (PCT Rule 17.2(a)).	on No ed in this National Stage			
Attachmen	et(s) te of References Cited (PTO-892)	4) ☐ Interview Summary	(PTO-413)			
2) Notic 3) Infor	the of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO/SB/08) or No(s)/Mail Date	Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ite			

RESPONSE TO APPLICANTS' AMENDMENT

Applicants' Amendments

Acknowledgment is made of Applicants' amendment filed 07/11/07, 05/16/07 and 01/12/07 in response to the non-final Office Action mailed 10/12/06. The former is compliant with 37 CFR 1.121. Applicants have amended the specification and the claims.

Status of Claims

2) Claims 8-13 and 38-51 have been canceled via the amendment filed 07/11/07.

Claims 1-7, 14, 19, 21, 22, 27, 29, 30, 35 and 37 have been amended via the amendment filed 07/11/07.

New claim 52 has been added via the amendment filed 07/11/07.

Claims 1-7, 14-37 and 52 are pending and are under examination.

Prior Citation of Title 35 Sections

3) The text of those sections of Title 35 U.S. Code not included in this action can be found in a prior Office Action.

Prior Citation of References

4) The references cited or used as prior art in support of one or more rejections in the instant Office Action and not included on an attached form PTO-892 or form PTO-1449 have been previously cited and made of record.

Objection(s) Withdrawn

- The objection to the specification made in paragraph 7(a) of the Office Action mailed 10/12/06 is withdrawn in light of Applicants' amendment to the specification.
- The objection to the specification made in paragraph 7(b) of the Office Action mailed 10/12/06 is withdrawn in light of Applicants' amendment to the specification.
- 7) The objection to the specification made in paragraph 7(c) of the Office Action mailed 10/12/06 is withdrawn in light of Applicants' amendment to the specification.

Objection(s) Maintained

8) The objection to claim 2 made in paragraph 7(c) of the Office Action mailed 10/12/06 is

maintained for reasons set forth therein and herein below. Although Applicants have amended a part of the specification to address the issue, several other trademark recitations continue to appear in the instant specification. For example, see 'Marcol 52' at line 28 of page 24. It is suggested that Applicants examine the whole specification and capitalize all trademark recitations, wherever such recitations appear.

Rejection(s) Moot

- 9) The rejection of claims 8 and 9 made in paragraph 9 of the Office Action mailed 10/12/06 under 35 U.S.C. § 101 as being directed to non-statutory subject matter, is most in light of Applicants' cancellation of the claims.
- 10) The rejection of claims 8 and 9 made in paragraph 11 of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is most in light of Applicants' cancellation of the claims.
- 11) The rejection of claims 8 and 9 made in paragraph 14 of the Office Action mailed 10/12/06 under 35 U.S.C § 102(b) as being anticipated by Quaissi *et al.* (*Science* 234: 603-606, 1986) as evidenced by Qin *et al.* (*J. Immunol.* 150: 2072-2080, 1993) and Haites *et al.* (*J. Biol. Chem.* 280: 10981-10987, 2005), is moot in light of Applicants' cancellation of the claims.

Rejection(s) Withdrawn

- 12) The rejection of claim 7 and those dependent therefrom made in paragraph 9 of the Office Action mailed 10/12/06 under 35 U.S.C. § 101 as being directed to non-statutory subject matter, is withdrawn in light of Applicants' amendment to the claim.
- 13) The rejection of claim 1 made in paragraph 11(a) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 14) The rejection of claim 1 made in paragraph 11(b) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 15) The rejection of claim 2 made in paragraph 11(c) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants'

Application/Control No. 10/749,143

Art Unit: 1645 September 2007

amendment to the claim.

- 16) The rejection of claim 3 made in paragraph 11(d) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 17) The rejection of claim 4 made in paragraph 11(e) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 18) The rejection of claim 5 made in paragraphs 11(f) and 11(g) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 19) The rejection of claim 6 made in paragraph 11(h) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- **20)** The rejection of claim 4 made in paragraph 11(j) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 21) The rejection of claim 19 made in paragraph 11(1) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 22) The rejection of claim 21 made in paragraph 11(m) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 23) The rejection of claim 27 made in paragraph 11(n) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- 24) The rejection of claim 29 made in paragraph 11(o) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.

Application/Control No. 10/749,143 Art Unit: 1645

September 2007

25) The rejection of claim 35 made in paragraph 11(p) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.

- 26) The rejection of claim 37 made in paragraph 11(q) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claim.
- The rejection of claims 2-7 and 14-37 made in paragraph 11(r) of the Office Action mailed 10/12/06 under 35 U.S.C. § 112, second paragraph, as being indefinite, is withdrawn in light of Applicants' amendment to the claims.
- 28) The rejection of claims 1-4, 14, 16, 18, 19, 22, 24, 26, 27, 30, 32, 34 and 35 made in paragraph 13 of the Office Action mailed 10/12/06 under 35 U.S.C. § 102(b) as being anticipated by Brodeur *et al.* (*Infect. Immun.* 50: 510-516, 1985) as evidenced by Rodriquez *et al.* (US 5,286,484), is withdrawn in light of Applicants' amendment to the claims and/or the base claim.
- 29) The rejection of claims 5-7 and 14-37 made in paragraph 14 of the Office Action mailed 10/12/06 under 35 U.S.C § 102(b) as being anticipated by Quaissi *et al.* (*Science* 234: 603-606, 1986) as evidenced by Qin *et al.* (*J. Immunol.* 150: 2072-2080, 1993) and Haites *et al.* (*J. Biol. Chem.* 280: 10981-10987, 2005), is withdrawn in light of Applicants' amendment to the claims.

New Rejection(s) Necessitated by Applicants' Amendment Rejection(s) under 35 U.S.C. § 112, First Paragraph (New Matter)

30) Claims 1 and those dependent therefrom are rejected under 35 U.S.C. § 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. This is a new matter rejection.

Claim 1, as amended, includes the new limitation: 'or a polypeptide at least 90% identical thereto and having a molecular weight of 44 kD to 55 kD as determined in SDS-PAGE, wherein the polypeptide is a polypeptide of *Neisseria meningitidis*'. The amendment to claim 1 deletes the previous limitation 'NMASP', and thereby broadens the scope of the claim. A generic polypeptide at least 90% identical to any isolated *N. meningitidis* polypeptide having a molecular weight of 44

kD to 55 kD as determined by SDS-PAGE, including a non-NMASP polypeptide, encompasses a truncated N. meningitidis polypeptide that has 10% of the amino acids deleted, i.e., a polypeptide that lacks 10% of the amino acids, and that still has a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE. However, there is no descriptive support in the instant specification for a Neisseria meningitidis polypeptide that is at least 90% identical to an isolated polypeptide having a molecular weight of 44 kD to 55 kD as determined in SDS-PAGE and having a molecular weight of 44 kD to 55 kD as determined in SDS-PAGE. Applicants state that the amendment is supported at least in paragraph 0046. However, there is no paragraph that is identified as paragraph 0046 in the instant specification, as originally filed. The original claim 1 and the first and second paragraphs under the 'Detailed Description of the Invention' are supportive of an isolated meningococcal NMASP having a molecular weight of about 40 kD to about 55 kD, preferably about 44 kD to about 53 kD as determined by SDS-PAGE. The currently claimed polypeptide however encompasses a non-NMASP polypeptide of Neisseria meningitidis having the recited molecular weight, for which there is no support. Furthermore, an isolated truncated polypeptide with 10% of the amino acids deleted would not be expected to have the same molecular weight as the nontruncated polypeptide from which it is derived. Therefore, the above-identified limitation in the instant claim(s) is considered to be new matter. In re Rasmussen, 650 F2d 1212 (CCPA, 1981). New matter includes not only the addition of wholly unsupported subject matter but also, adding specific percentages or compounds after a broader original disclosure, or even omission of a step from a method. See M.P.E.P 608.04 to 608.04(c).

Applicants are respectfully requested to point to the descriptive support in the specification as filed by pointing to specific lines and pages, for the new limitations, or alternatively, remove the new matter from the claim(s). Applicants should specifically point out the support for any amendments made to the disclosure. See MPEP 714.02 and 2163.06.

31) Claim 7 and dependent claims 15, 17, 20, 21, 23, 25, 28, 29, 31, 33, 36 and 37 are rejected under 35 U.S.C. § 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. This is a new matter rejection.

Application/Control No. 10/749,143

Art Unit: 1645 September 2007

Claim 7, as amended, is drawn to an isolated peptide 'consisting of an immunogenic fragment of 7 or more amino acids of a polypeptide of Neisseria meningitidis having a molecular weight of '44 kD to 55 kD' as determined by SDS-PAGE'. The amendment to claim 7 deletes the previous limitation 'NMASP', and thereby broadens the scope of the claim. As currently claimed, the peptide encompasses a peptide consisting of an immunogenic fragment of 7 or more amino acids of a non-NMASP polypeptide of N. meningitidis having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE. Applicants state that support for the limitation '7 or more' can be found in the first paragraph under Section 5.2 on page 13. However, this part of the disclosure is limited to a fragment of the 'NMASP' polypeptide of N. meningitidis having a molecular weight of 44 kD to 55 kD or 44 to 53 kD as determined by SDS-PAGE wherein the fragment has 7 or more amino acids of said NMASP. This part of the specification does not identify the peptide 'consisting' of seven amino acids of the NMASP polypeptide, let alone any other polypeptide of a molecular weight of 44 kD to 55 kD, to be 'immunogenic'. Therefore, the above-identified limitation in the instant claim(s) is considered to be new matter. In re Rasmussen, 650 F2d 1212 (CCPA, 1981). New matter includes not only the addition of wholly unsupported subject matter but also, adding specific percentages or compounds after a broader original disclosure, or even omission of a step from a method. See M.P.E.P 608.04 to 608.04(c).

Applicants are respectfully requested to point to the descriptive support in the specification as filed by pointing to specific lines and pages, for the new limitations, or alternatively, remove the new matter from the claim(s). Applicants should specifically point out the support for any amendments made to the disclosure. See MPEP 714.02 and 2163.06.

Rejection(s) under 35 U.S.C. § 112, First Paragraph

32) Claim 1 and those dependent therefrom are rejected under 35 U.S.C. § 112, first paragraph, as based on a disclosure which is not enabling. Claim 1 is not enabled because the sequence identifying (SEQ ID) number for the isolated *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE, critical or essential to the practice of the invention, is not included in the claim(s). *In re Mayhew*, 527 F.2d 1229, 188 USPQ 356 (CCPA 1976).

The feature of being 'at least 90% identical' to an isolated *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE is essential for the

Application/Control No. 10/749,143

Art Unit: 1645 September 2007

claimed polypeptide. However, the SEQ ID number of the isolated *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE is not recited in claim 1. As illustrated under art rejections(s) previously and herein below, *N. meningitidis* produces several polypeptides having a molecular weight in the range of 44 kD to 55 kD as determined by SDS-PAGE, the precise structure of all of which is not known. The claim lacks the critical limitation with regard to the amino acid sequence of *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE, without which one of ordinary skill in the art cannot perform the structural comparison, and practice the invention as currently claimed. The instant claim does not structurally identify the recited *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE by a sequence ID number. A polypeptide having at least 90% identity can be obtained only when the sequence(s) to be compared is identified in the claim(s) by a specific structure or SEQ ID number. Without such a disclosure, undue experimentation would have been required by one of ordinary skill in the art to practice the invention as claimed.

33) Claim 7 and dependent claims 15, 17, 20, 21, 23, 25, 28, 29, 31, 33, 36 and 37 are rejected under 35 U.S.C. § 112, first paragraph, as based on a disclosure which is not enabling. Claim 7 is not enabled because the sequence identifying number for the recited *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE, critical or essential to the practice of the invention, is not included in the claim(s). *In re Mayhew*, 527 F.2d 1229, 188 USPQ 356 (CCPA 1976).

The feature of 'consisting' of an immunogenic fragment of '7 or more amino acids' of a *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE is essential for the claimed peptide product. However, the SEQ ID number of the recited *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE is not recited in the claim. *N. meningitidis* produces several polypeptides having a molecular weight in the range of 44 kD to 55 kD as determined by SDS-PAGE, the precise structure of all of which is not known. The claim lacks the critical limitation with regard to the amino acid sequence of *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE, without which one of ordinary skill in the art cannot identify and obtain a 7 or more amino

acid-long peptide as currently claimed. The instant claim does not structurally identify the recited *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE by a specific sequence ID number. A peptide consisting of an immunogenic fragment of 7 or more amino acids as recited can be obtained only when the sequence(s) from which it is obtained is recited by identifying the polypeptide with a SEQ ID number. Without such a disclosure, undue experimentation would have been required by one of ordinary skill in the art to practice the invention as claimed.

Rejection(s) under 35 U.S.C. § 112, Second Paragraph

- 34) Claims 1-7, 14-37 and 52 are rejected under 35 U.S.C. § 112, second paragraph, as being indefinite, for failing to particularly point out and distinctly claim the subject matter which Applicants regard as the invention.
- (a) Claim 1, as amended, is indefinite and confusing in the limitation: 'a polypeptide at least 90% identical thereto and having a molecular weight of 44 kD to 55 kD as determined in SDS-PAGE'. It is unclear how a polypeptide having at least 10% non-identity to a 44 kD to 55 kD polypeptide can have a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE. Does it mean that a polypeptide having at least 10% of the amino acid residues deleted from a 44 kD to 55 kD polypeptide would also have a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE?
- (b) Claim 1, as amended, is indefinite in the limitation: 'a polypeptide at least 90% identical thereto', because it is unclear what is encompassed in this limitation. Since there is no SEQ ID number recited in the claim, it is unclear whether 90% identity represents 90% functional identity.
- (c) Claim 1 is vague and indefinite in the limitation 'a polypeptide at least 90% identical thereto'. The claim neither identifies the amino acid sequence of the isolated *N. meningitidis* polypeptide having a molecular weight of 44 kD to 55 kD as determined by SDS-PAGE, nor the amino acid sequence of a polypeptide at least 90% identical thereto for one of skill in the art to perform the comparison, or to envisage the metes and bounds of the claim. It is unclear how the percent identity of at least 90% is determined, i.e., in comparison to what other SEQ ID number. The only comparable structural limitation of the polypeptide recited in the instant claim is the

molecular weight of the polypeptide. However, one cannot use molecular weight to obtain a certain percent identity.

- (d) Claim 3, as amended, is vague, indefinite and/or confusing in the limitation: 'serogroup Types A-L and W', because it is unclear what is encompassed in the limitation 'serogroup Types A-L and W'. Do the 'serogroup Types' represent *Neisseria meningitidis* serotypes, immunotypes, subtypes, or serogroups?
- (e) Analogous rejection and criticism apply to claim 4, as amended, with regard to the limitation: 'serogroup Type A, Type B, Type C or Type W'.
- (f) Claim 6 is indefinite and confusing in the Markush language: 'selected from the group consisting of SEQ ID NO: 11'. Since there is only one Markush species, it is suggested that Applicants replace the above-identified limitation with the limitation --of SEQ ID NO: 11--.
- (g) Claims 2-6, 14-37 and 52, which depend directly or indirectly from claim 1 or claim 7, are also rejected as being indefinite because of the indefiniteness identified above in the base claim.

Rejection(s) under 35 U.S.C. § 102

35) Claims 1-4, 14, 16, 18, 19, 22, 24, 26-30, 32 and 34-37 are rejected under 35 U.S.C § 102(b) as being anticipated by Allunans *et al.* (*APMIS* 106: 1181-1187, December 1998) as evidenced by Schlecht *et al.* (*Naturwissenschaften* 80: 9-17, 1993, abstract).

Instant claims are granted the effective filing date of the instant application due to the new matter identified above, and therefore Allunans *et al.* qualifies as prior art under 35 U.S.C § 102(b).

The limitation 'vaccine' in claims 22, 24 and 26-29 represents the intended use of the claimed polypeptide composition and therefore is not given any patentable weight in this rejection.

Allunans *et al.* taught an isolated or purified polypeptide of a serogroup (i.e., serogroup Type) A *Neisseria meningitidis* having a molecular weight of 47-48 kDa as determined in SDS polyacrylamide gel electrophoresis and contained in a phosphate buffer. See abstract; Figure 3; paragraph bridging pages 1185 and 1186; paragraph bridging two columns on page 1183; Materials and Methods; and section 'SDS-polyacrylamide electrophoresis on page 1183. The molecular weight of the prior art polypeptide falls well within the recited range of '44 kD to 55 kD' and '44 to 53 kD' and therefore, the prior art isolated or purified polypeptide anticipates the instantly claimed

polypeptide. Since the prior art polypeptide is not fully purified, it is expected to contain at least residual or contaminant phospholipids, or cell wall components such as OMP, lipoprotein, and the meningococcal lipopolysaccharide immunogen. Because of the overlapping molecular weight and the bacterial origin of the prior art polypeptide, the prior art polypeptide is viewed as the same as the polypeptide claimed in the instant claims, and therefore it is expected to have the same intrinsic structure and properties as that of the Applicants' polypeptide. That one or more of the residual cell wall components such as OMP, lipoprotein, and the meningococcal lipopolysaccharide in the prior art polypeptide serve as intrinsic immunoadjuvants is inherent from the teachings of Allunans *et al.* in light of what is well known in the art. For example, Schlecht *et al.* taught that bacterial cell wall components such as lipopolysaccharides, a variety of membrane proteins, murein, and lipoprotein act as immunoadjuvants. See abstract of Schlecht *et al.*

Claims 1-4, 14, 16, 18, 19, 22, 24, 26-30, 32 and 34-37 are anticipated by Allunans *et al*. The reference of Schlecht *et al*. is not used as a secondary reference in combination with the reference of Allunans *et al*., but rather is used to show that every element of the claimed subject matter is disclosed by Allunans *et al*. with the unrecited limitation(s) being inherent as evidenced by the state of the art. See *In re Samour* 197 USPQ 1 (CCPA 1978).

36) Claims 1-7, 14-37 and 52 are rejected under 35 U.S.C § 102(b) as being anticipated by Jackson *et al.* (WO 00/12535, published 03/09/2000) ('535).

Instant claims are granted the effective filing date of the instant application due to the new matter identified above, and therefore Jackson et al. qualifies as prior art under 35 U.S.C § 102(b).

Jackson et al. ('535) disclosed an isolated NMASP polypeptide of Neisseria meningitidis having a molecular weight of 44 kD to 55 kD, or 44 kD to 53 kD as determined by SDS-PAGE wherein the polypeptide comprises an amino acid sequence that is 100% identical to SEQ ID NO: 11 that is recited in the instant claim 52. Jackson et al. ('535) also disclosed the seven amino acidlong peptide, GNSGGPL, of said NMASP polypeptide of Neisseria meningitidis. See page 20; and claims 7-9. A vaccine or an antigenic composition comprising the polypeptide or the peptide, with or without a pharmaceutically acceptable carrier, one or more adjuvants, lipids, lipopolysaccharides, proteins, phospholipids, attenuated organisms, and/or inactivated whole cells is taught. The polypeptide is from Neisseria meningitidis Types A, B, C, or W, and binds to an antibody that

specifically binds to a protein having the amino acid sequence of SEQ ID NO: 11. See abstract; pages 70 and 71; and claims 5, 6, 1-4, and 14-38.

Claims 1-7, 14-37 and 52 are anticipated by Jackson et al. ('535).

Remarks

37) Claims 1-7, 14-37 and 52 stand rejected.

To be consistent with the claim language used in claim 7, it is suggested that Applicants replace the limitation 'as determined in ...SDS-PAGE' in claim 1 with the limitation --as determined by SDS-PAGE--.

38) Applicants' amendment necessitated the new ground(s) of rejection presented in this Office action. THIS ACTION IS MADE FINAL. Applicants are reminded of the extension of time policy as set forth in 37 C.F.R 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 C.F.R 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

- **39)** Papers related to this application may be submitted to Group 1600, AU 1645 by facsimile transmission. The Fax number for submission of amendments, responses and/or papers is (571) 273-8300, which receives transmissions 24 hours a day and 7 days a week.
- 40) Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAG or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.Mov. Should you have questions on access to the Private PAA system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-

Application/Control No. 10/749,143 Art Unit: 1645

September 2007

9199 (in USA or CANADA) or 571-272-1000.

41) Any inquiry concerning this communication or earlier communications from the Examiner should be directed to S. Devi, Ph.D., whose telephone number is (571) 272-0854. A message may be left on the Examiner's voice mail system. The Examiner can normally be reached on Monday to Friday from 7.15 a.m. to 4.15 p.m. except one day each bi-week, which would be disclosed on the Examiner's voice mail system.

If attempts to reach the Examiner by telephone are unsuccessful, the Examiner's supervisor, Jeffrey Siew, can be reached on (571) 272-0787.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (571) 272-1600.

September 2007

